I was happy when Mrs. Gimbel and Dr. Rinkel invited me to come to participate in this meeting, and I have been happy to attend the symposium on the chemical basis of mental disease and to be associated with the Manfred Sakal Foundation, because of its dedication to the job of diminishing the amount of human suffering in the world.

Recently, the world has been changing at a tremendous rate. Technological advances are such that the world is not the same one year as it was the year before. Tremendous weapons of destructions are developed. The stresses under mi which people live become ever more oppressing. There emanates from the centers of governments a special sort of irrationality. There are rejections of dogma/ and revelation, the old principles of conduct. We are forced to strive for a fundamental, ethical, and philosophical principle under which we can live, if the world is going to survive.

I think that the fundamental ethical principle that everyone can accept is that of the minimization of human suffering. This is not the same as maximizing the amount of happiness in the world. If we take money as the criterion, to increase the income of, say, a billion people by a hundred dollars a year is not at all equivalent in the amount of happiness that it causes to decreasing the income of the other two billion by half of that amount. It is the suffering, human suffering that we have to pay attention to.

As the means of communication and transportation, telephones, telegraphs, the press have developed in the world, the nature of the human race has changed. No longer are individual human beings the units. They are now bound together by these means of communication in the same way that the cells of an individual human body are bound together by the nerves that interconnect them, in such a way that the whole human race is becoming one organism, and a special ethical

be an ethical principle that places the survival of society or the race above the value, the worth of the individual human being. It is the individual human being and his suffering to whom we must pay attention:

Now there are many causes of human suffering. In some parts of the world starvation, malmutrition, infectious diseases are the principal causes; but everywhere mental disease is an important cause, and the half of the hospital beds in the country are occupied by mental patients. Mental disease causes a particularly great amount of suffering for two reasons:

First, it may continue in an individual year after year, it fluctuates causing him great suffering for many and his friends and relatives. And second, often it attacks some of the that most able of our people, also start interfactual lines.

It may well be that schizophrenia is an especially significant disease for the most able intellectuals. This is a disease that we must attack. We must cut down on the amount of suffering caused by mental illness.

I am grateful to the psychiatrists. For several years helf a dezen wears. I have had close contact with them, net, as some people have suggested because I hereafted professional service, has because I became interested the in the question of how what chemistry might contribute to medicine, might be used in diminishing the amount of human suffering, and decided that mental disease was the field in which there was the greatest need of effort.

poor job because they don't know how to do any better. They use crude methods, insulin shock, electro-shock therapy, damaging the whole human organism in the hope that the new changed individual will be in some way improved.

These drugs were not discovered with the use drugs attraction of the standard of the diseases; Inc. horses by purely empirical processes, it has been found that for some patients, these days are helpful, reliable.

I hope that the psychiatrists will not have to work longer under this terrible handicap of extreme ignorance, and I am encouraged by the interesting presented at this symposium information about most interesting valueble land about abnormalities in the biochemistry of schizophrenia individuals or other people who are suffering from mentalized disease, as compared with other human beings.

I am encouraged to believe that the time will come before long, in ten twenty years when, if enough attention is paid to this jeb, there will be some significant fundamental, basic understanding of the nature of the mental diseases, comparable to that that exists now for a few other diseases that are called molecular diseases.

I believe that most mental diseases are molecular diseases, the result of a biochemical abnormality in the human body. I think that the psyche, that consciousness, the mind, is a manifestation of the structure of the brain, an electrical oscillation in the brain supported by the material structure of the brain comblement being altered by an abnormality made abnormal by an abnormality in the material structure of the brain itself, usually second with the property of the brain itself, usually second by an abnormality in the environment.

Pallagra is an example of a disease the responsible, that had a mental manifestations, was prevalent years ago. It was a deficiency disease, due to efficiency disease, due to constant a lack of vitamin, nicotinic acid. This discovery has led to the solution of the pallagra problem in many parts of the world.

phenylketonuria, first identified 30 years ago by Dr. Feeling in Oslo, Norway, who checked up on some mentally deficient children who had an odd smell and found that there were some unusual substances present in their urine; and the further investigation by him and other scientists led to a discovery that these children lack an enzyme in the liver that catalages the oxidation of an amino acid, phenylalanine to tyrosine, so that the phenylalanine builds in the bloodstream and cerebrospinal fluid to high concentrations where the total interferes with the development of and function of the brain and leads to mental deficiency.

Foelling

A Property of the Parketing Pauling 5 One person in 80 CE Penson Carries A gene people have two genes that manufacture the enzyme in the in liver that catalyzes the wridation of presynaterness, You know, the proteins that we eat such as in the meet and the fish and potatoes and bread the proteins that we tak contain about 20 different acids: glyscine, alanine, derine, tyresine, melalinicine and so on. We eat more than we need of it not quite that enough tyresine. There is an in the liver that enough tyresine, the enzyme of catalyzes the oxidation of Most people have two genes that they have inherited from their parents, that manufacture the enzyme in the liver to do this job. One person in 80 has one gene that manufactures the enzyme, and one damaged, mutated appropriate gene that went, manudo the job or else there to all abnormal to ayen molecula that will not do its the have only half the amount enzyme that a normal person has. But that's enough to keep

them in good health.

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tody fluids and interfered with the growth and functioning

But if two of these people marry one another, that there occurs the great lottery; the the child inherits of every pair of genes from the finder and the mether of the two genes from the father one of the two from the mother. A quarter of the children will be inherit at abnormal gene of the father and an abnormal gene of the mother. Weither will manufacture the enzyme.

There is none of this enzyme in the liver. As the child eats its food the phenylalanine builds up a second causing him to be mentally deficient.

But when this knowledge was obtained, it was recommend that is not the first enzyme of the second of the manifestations of the manifestations of the second of the manifestations of the second of the manifestations of the manifestation of the manifestation of the manifestation of the manifestation of the manifestation

But when this knowledge was obtained, it was recommend that is a should test for this effect at the standard a few weeks after birth. You should test for this effect at the hydrolyzed protein bottom of the standard from which the max phenylalanine had been removed at this paper at the time, instead of ordinary protein.

of the institutionalized mentally deficient individuals in the United States, now may be brought under control because of the discovery about its nature.

It is not one that leads to mental deficiency, but it is molecular disease, with about which I know more than any me other, and about which I have a personal connection, because its molecular nature was discovered by three of my students working in Pasadena 13 years ago.

It is a disease of the blood, and a very important one. Perhaps

10 or 20 thousand people in the world die of it. Children are born with this

disease; they suffer for a few years and then die the suffer some of the suffer for a few years.

to be 20 person old, persons. Calle is caused, tan, by such a billion.

twisted out of shape. Often, instead of being flat, they are twisted into a sort of crescent shape that looked to Dr. Herrick in 1911 sately like a sinkesickle with which one cuts grass. He named is lickle cell amenia. It is easier to study the blood than the brain. It is easier to study hemselbline than the brain, say: The brain is going to be a tough mut to erack for people who are working on mental disease.

human being, makes up to 1 per cent of the weight of an interest of human being. The season can get quite a bit out of anybody if you want in investigation. It is a make beautiful protain, red, because the molecule, which has about 10,000 atoms in the has 4 ion atoms which are able to combine with oxygen in the lungs and carry the oxygen out to the extremities.

The red cells of these patients with Sickle cell anemia have these red cells which are twisted out of shape. They are only twisted out of place in the venous blood, not in the arterial blood. There they have the normal shape. As you see, there is the cleavage.

It is highly probable that it is the hemoglobin is molecule then that is responsible for the disease, because in arterial blood you don't have hemoglobin; you have instead oxygen; whereas in venous blood, there is something different. If these patients manufacture a sort of hemoglobin was different from that which normal individuals manufacture, these might be sticky molecules, self-complementary, such that they would clamp onto one another to form long rods which would line up side by side to form a long needle-like thing. It would be longer than the diameter of the red cell, and it would twist itself out of shape. They would be sticky and get tangled up with themselves and prevent the flow of blood through the capillaries and the various manifestitions

of the disease would occur.

So, Dr. Tunneau and Dr. Singer, and Dr. Wells, after a while, with encouragement, carried out an experiment. They put a little drop of make salt water, with electrodes at the ends -- a positive electrode and a negative electrode -- and introduced a drop of hemoglobin from a normal human being. In this colorless liquid, under the influence of the electric current, it moved over toward the positively-charged electrode. It had a negative electric charge.

When a drop firm of blood from a patient with the disease was put in the trough, it moved over to the negatively-charged; if it had a positive charge, it had to be different from the normal hemoglobin in the red cells of normal individuals.

If you mixed these two and put a drop of the mixture in the electric field, they separated; part of the blood moved this way; the rest that way. Then when they got blood from the father of the patient and the mother of the patient -- and got Dr. Tunneau of our laboratory put a drop of hemoglobin from the father of the patient -- that blood split in half; half moved toward the anode, and half toward the cathode; and similarly for the mother.

This made the nature of the disease clear. The father and the mother had one normal gene that manufactured normal hemoglobin. Each had a normal gene, and each had an abnormal gene, and each of these genes manufactured its own kind of hemoglobin. Here we had an abnormal hemoglobin. And the patients had inherited only the abnormal gene of the father and only the abnormal gene of the mother.

Here was the demonstration, for the first time, of a disease of the hemoglobin molecule that produced the manifestations of the disease Sickle

cell anemia.

That isn't the whole of the story. A month or two later, Dr. Tunneau brought in some blood from another patient who had a still more abnormal hemoglobin, which was named Hemoglobin C, and then Hemoglobin D, Hemoglobin E, Hemoglobin G, Hemoglobin H, Hemoglobin I -- some 30 or 40 of these. I haven't kept track of them. Otu of these, abnormal hemoglobins have been found, many of them associated with the disease, even with diseases we that result in sort of hybrid diseases that result from the inheritance of a Sickle cell amenia from one parent and the inheritance of a C gene from the other. Interespense Neither of these genes alone, in single dose, causes a disease that amounts to anytying, but the patient who has inherited both the first abnormality and the second abnormality in single dose has a disease, a new kind of hybrid disease, hemolytic anemia.

This example, I think, indicates what we can expect to find in schizophrenia, not that there is a gene for schizophrenia, such that when the patient inherits this gene in Souble dose he is schizophrenic, but rather that there are many genes such that any one of them, when present in double dose, or perhaps one single dose, and another one in single dose, or it may even be a combination of several of these genes, produces an abnormality, a quantitative abnormality to make it difficult for the individual to accept reality, to think in the normal state, or to retain his sanity under the normal stresses of living, or increasing stresses of living in the modern world.

There may be factors involved in schizophrenia that come from the outside deficiency factors, some lack of some vitamins, or perhaps toxic chemical substances that are present. Who knows?

The evidence that was presented to the symposium about the presence of some sort of a biochemical abnormality in the blood associated in some way with

the globulin fraction of the serum of the blood seem to me to be very suggestively convincing, because several groups of good investigators from different places using different methods of investigation had obtained somewhat similar results with the methods.

I feel this is a strong indication of an abnormality that is present also in the brain and that is involved in the disease -- in some of the diseases that we classify as schizophrenia. Of course, this multiple character of a disease will make it difficult to find a cure, a treatment. But for all kinds of schizophrenia, I can't say that I have any hope that mental disease can be brought completely under control. But I do believe that a tremendous amount of progress can be made in controlling it, and in decreasing the amount of human suffering that is involved.

There are ways for treating diseases that can be envisaged that have not yet been brought into practice and that are worth serious effort. For example, in an enzyme deficiency disease, galactocemia, or many of the other enzyme deficiency diseases. With the progress in our knowledge about the nature of enzymes, the structure of protein molecules, it will before long be possible to synthesize artificial enzymes that will have enzymic activity to perhaps implant a capsule containing some stable artificial enzyme as a substitution therapy that permits the patient to get along well in the course of his life, possibly with an occasional replacement.

New drugs will be discovered, some of them by the same max emperical process, by means of which the Aborigines discovered the 1 efficacy of chewing tobacco leaves in the treatment of malaria.

But the time will come when we shall have enough understanding of the nature of mental disease to be able to synthesize drugs to order that will be specific for a particular disease that will operate not in the rather generally nonspecific manner of the present drugs that are effective, that have changed

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the character of mental hospitals greatly, but that nevertheless will still have to be described as representing a sort of shotgun attack on the problem of the solution of disease.

In the case of diseases, congenital diseases involving gene abnormalities, \mathcal{N}^{h} we have now to recognize the possibility of the Das replacement -- replacement introduction into the defective individual of \mathcal{D}^{hh} molecules that will serve the purpose of the molecules that they did not inherit from their parents.

But, all of these are palliative measures. They are not the solution of the problem. The pool of human germ plasma, you know, continues to degenerate day after day, as the natural mutagenic agents, natural radioactivity, other mutagenic substances, and those chemicals and artificial high energy radiation that constitute a part of our modern life damage the Dea molecules constituting the pool of plasma. It is a very precious one.

A person can inherit about 100,000 molecules of Dan, 50,000 from his father, and 50,000 from his mother. I all of these inherited by every one of the 2 billion persons now living on earth were to be brought together -- could be brought together -- they would constitute a mass of about 4 milligrams, the size of 4 pinheads.

This is the pool of human germ plasma that I am concerned about. It becomes damaged -- damaged day after day by mutagenic agents, and it is purified. There is a natural process of purification that goes on. New genes for phaserix phenyluria are caused by this mutation process. The probability is something like one in a hundred births, or one in 50,000 children has a new gene for this. For example, there is a dominant gene that causes dwarfism. It is a dominant gene. The parents of a child who did not possess the abnormal genes, but had two normal genes instead, would between the conception of the parents and the conception of the child produce an abnormal gene that he inherited.

One child in 12,000 inherits this.

There is the disease cystic fibrosis. One child in 800 has inherited from its parents two genes for cystic fibrosis which give him this very serious disease -- and we don't know what these genes do. This is without doubt a molecular disease, but the protein molecule that is abnormal has not been identified.

Now when these defective children are born and die without progeny, they remove from the pool of human germ plasma 2 of the defective genes, and so there has been a steady state, such that some 4 per cent of the children who are born have some degree of mental defect; some 7 per cent are congenitally defective k either physically or chemically.

I believe that it is possible for us now to begin to carry out the process of purifying the pool of human germ plasma at a rate to keep up with this continued contamination, without the suffering that is involved in the birth and death, # without progeny of the defective children. If two parents have a phenylalanine, then it is known that they are carriers of the gene for phryaphenylketomuria for each successive child; the chances are 25 per cent that the child, this child, too, will be defective. If two parents have a child with cystic fibrosis, or with many other diseases, Tasacks disease, nimintax disease, we developed a test, a rather difficult one, for detecting the heterozygous for phenylketonuria. One of the parents involved came to see us later. His wife had died. He knew that he and his first wife, if they had had further children, there would have been the 25 per cent chance. He came with a young woman that he wanted to marry, and he said he wanted to know whether she carried the gene for phenylketonuria. After the test was made, he was told she was not a carrier for phenylketonuria; that they could get married, and that none of their children would have the disease phenylketonuria; of course they would have some other defects, because everyone inherits some congenital defects, mainly minor ones. The minor ones are not to be laughed at. They are a cause of a tremendous amount of human suffering, because they are not enough to prevent procreation, and the genes are passed on for generation after generation causing continuous adding to the amount of human suffering in the world.

But it is possible in many cases now to say that affairning offspring in this marriage would have a 25 per cent, each child being grossly defective. This is a great amount of human suffering, and I would say that it is too great a chance to be taken; that a chance as great as 25 per cent of giving birth to a defective child, increasing the amount of human suffering is so great that this matter should not be left to a combination of ignorance combined with free enterprise in life as is customary at the present time.

People need knowledge. People need to know what their situation is, and then of course to decide for themselves in these personal matters. I do not advocate legal action. I do not advocate eugenics as a means of improving the race. We don't know enough about how to improve the race; but we do know something about how to decrease the amount of human suffering, and if a heterozyal were to marry a normal individual as this man, the father did, then half of those children would inherit his gene. If the only thing that was done as a result of minx increase in knowledge was to get people who possessed defective, seriously defective genes of a certain sort to marry other people who do not possess these genes, then there would be no elimination of the defective genes from the pool of human germ plasma, and pretty soon everyone would have a greater chance of being defective than he has at the present time.

I think that in the case of people who know that they are carrying a gene, a recessive gene for a serious disease, should marry normal individuals, and then have a somewhat smaller number of children than normal,

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one or two children, rather than three or four. In this way these defective genes would be removed from the pool of human germ plasma slowly, perhaps at a rate that they are now being removed by the death and suffering of the defective children, but without the death and suffering of these children. This, I believe, is a humane and rational way of attacking the problem of disease, congenital disease caused by mutated genes, a way that does not involve the birth and suffering of defective children, and as the years go by, we shall be able to obtain for mental diseases, too, more and more knowledge about the diseases and methods of predicting whether one union has a p high probability of leading to these defective births.

But, of course, we must also consider the question of palliative treatthat
ment. I wish textest the support for investigators who are studying the biochemical abnormalities of schizophrenic individuals could be doubled, tripled,
quadrupled. You know there are plenty of scientists, very able people in the
world, who would like to be working on problems like this, but who are instead
working on other problems the nature of which I haven't time to go into. But
we have no need to have technological scientific unemployment resulting from
disarmament. There is plenty of room for the scientist to do for the benefit
of human beings in the United States and all over the world.

I believe that we are going to have a better and a better world in the future, and that the scientists are going to do their part in contributing to it -- and of course with the help of the psychiatrists, people who work in the field of mental disease. We couldn't get anywhere without the help of the psychiatrists.

I want to say again that as I have become acquainted with more and more psychiatrists— and our eldest son is a psychiatrist now, 37 years old, and I think making a success in this profession, he seems to me to be a very good man, one of the soundest, most well-balanced men that I know. I think that

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he is good for his patients; no doubt does the best that he can for his patients. I hope that my associates and I out in Pasadema will, as we continue with our work, which is on a small scale, carried out not with the hope that we can make any mask epoch-making discoveries, but with the firm confiction that we can do something that will be of aid to him in his future work, and to all of the fine medical men in the world who are devoting themselves to the treatment of the mentally ill.

Thank you.